Amendments to the Specification:

Please replace the paragraph beginning at page 11, line 21, with the following amended paragraph:

"The commercial split flu antigen vaccine (Fluzone®) as used herein and prepared following the above procedure was obtained from Connaught Laboratories Inc., Swiftwater, PA, USA."

Please replace the paragraph beginning at page 17, line 19, with the following amended paragraph:

"Mice were bled one day prior to the first immunization and also on days 22 and 28 of the study. Immunizations were done on days 1 and 22. Both immunizations were administered intramuscularly in the thigh muscle. Each immunization was done at two injection sites (both right and left thigh muscles; 0.05 ml/site). The dose of RSV vaccine was 1 μg total protein and the dose of Fluzone® (split flu antigen) vaccine was 5 μg total protein per dose. The RSV or Fluzone® (split flu antigen) vaccines were administered in the presence or absence of adjuvant. The adjuvant used was poly-di(carboxylatophenoxy)-phosphazene (PCPP) given at 200 μg/dose. Mice that received live RSV (A2 strain) as the immunogen were given 1.5 x 10^6 pfu/dose intranasally. Mice that received live influenza virus (A/Taiwan Strain) as the immunogen were given 200 to 400 HAU/dose intraperitonally. Virus challenge with either RSV or influenza was administered intranasally on day 29 using the same dose as given for the live virus immunized mice. All animals were sacrificed on day 33. Lungs were removed and frozen immediately in liquid nitrogen for later determination of virus titre."